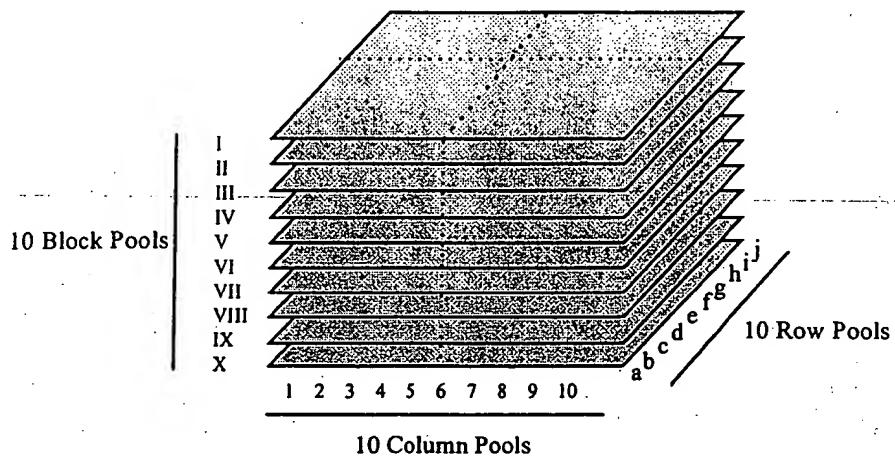
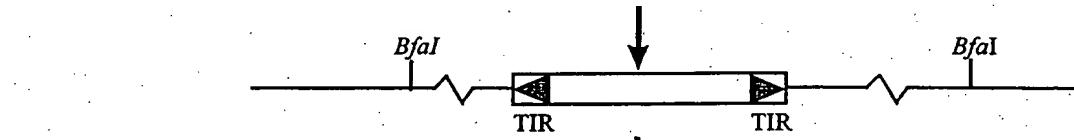
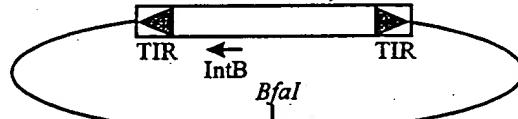
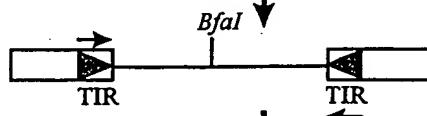


A: Amplification of pools of *dTph1* Flanking sequences

Harvesting of the 30 DNA pools of the 3D library

*BfaI* digest of the genomic DNA
Backligation of the fragmentsiPCR with IntA and IntB primer to
generate 30 pools of flanking sequences

Reamplification with nested IR primer

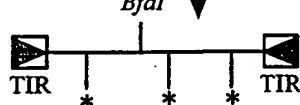


Figure 1A

B: Simple screening for insertion mutants

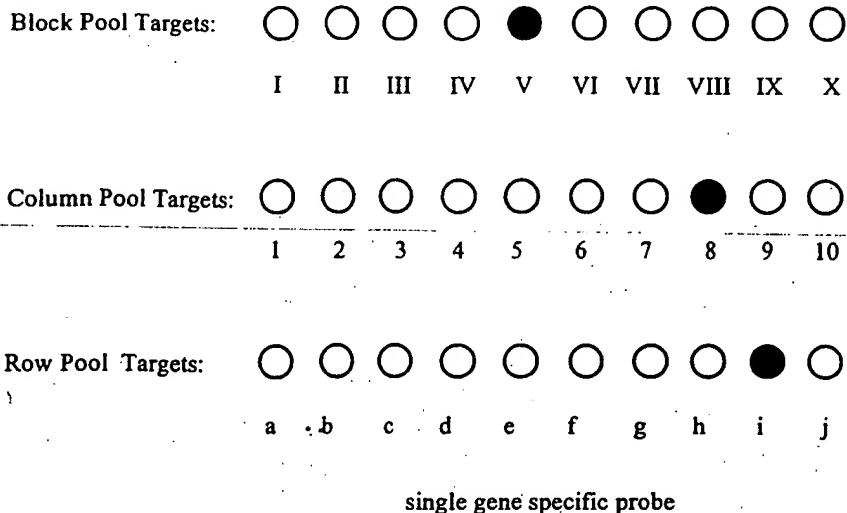


Figure 1B

C: Parallel screening for insertion mutants

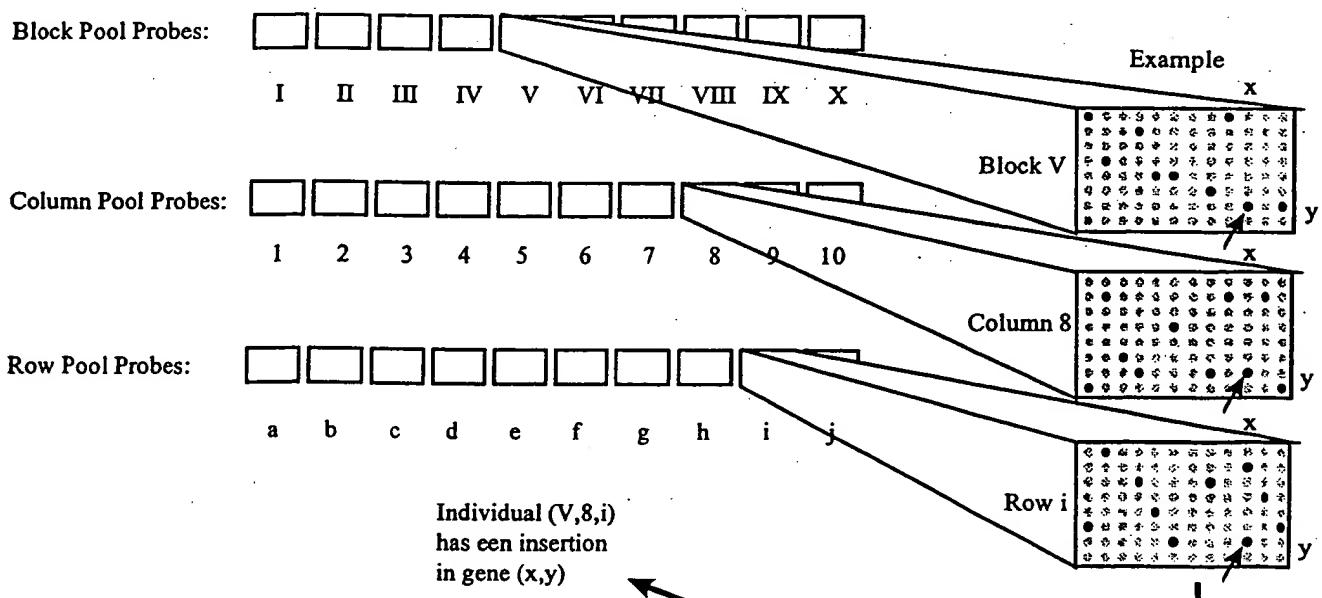


Figure 1c

Figure 1: Parallel screening for insertion mutants using iPCR: Schematic outline of the technique

A: Amplification of the *dTph1* flanking sequences: 1000 plants are organized in a 3D array, and 30 genomic DNA pools are prepared, each representing 100 individual plants. The 30 pools of genomic DNA are digested with a tetracutter enzyme (*Bfa*I). The genomic fragments are self-ligated and the insertion flanking sequences are amplified by PCR using two internal insertion specific primers, IntA and IntB. The specificity of the amplified products can be increased by nested reamplification of the flanking sequences using the terminal inverted repeat primer (TIR).

B: Simple screening for insertion mutants: The amplified flanking sequence pools are displayed on a filter and hybridized with a probe for any target gene of interest in a serial manner.

C: Parallel screening for insertion mutants: The 30 amplified flanking sequence pools are labelled and used to probe an array of target genes displayed on 30 replica filters (or microarrays) in a set of 30 hybridization reactions

**AMPLIFICATION and DETECTION of TRANSPOSON FLANKING SEQUENCES
USING TRANSPOSON DISPLAY (After Van den Broeck *et al.*, 1998)**

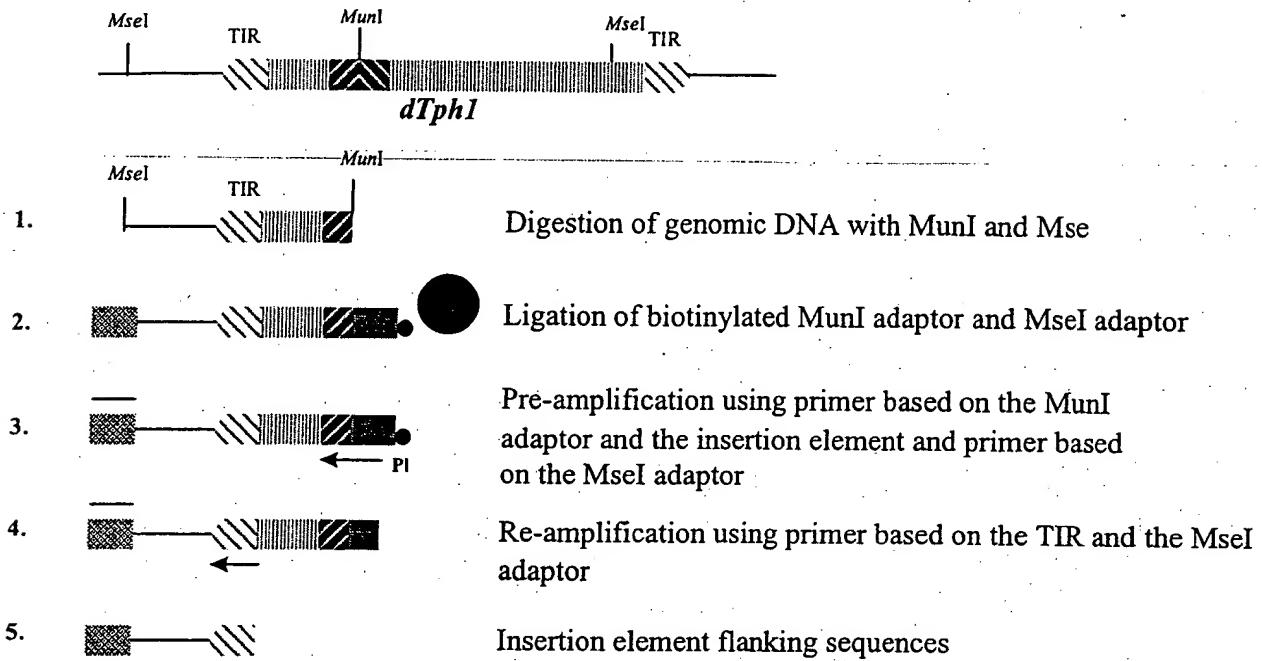


FIGURE 2

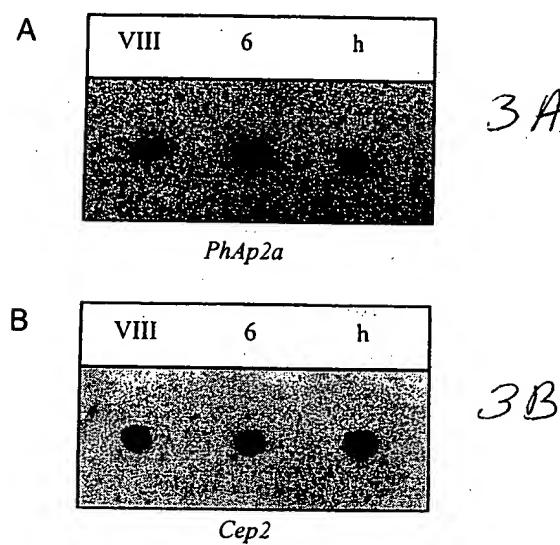


Figure 3: Generation of a 3D library of flanking sequences

DNA of an *PhAp2A* insertion mutant was added as a positive control to all the 3D pools, *dTph1* flanking sequences were amplified as described. Pools VII, 6, h were spot blotted and hybridized with A: a *PhAp2A* probe; B: a *Cep 2* probe.

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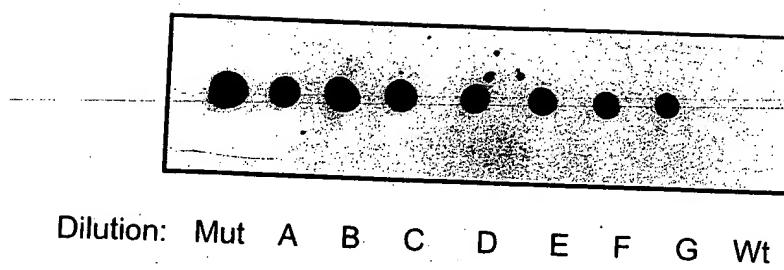


Figure 4a: 3D Screening with a 4Kb XbaI *Ap2A* fragment as a probe. Transposon display fragments spotted; A-G represent a dilution series as described in the text; Wt is wildtype. The probe was fluorescein-labeled, exposure time 5 minutes.

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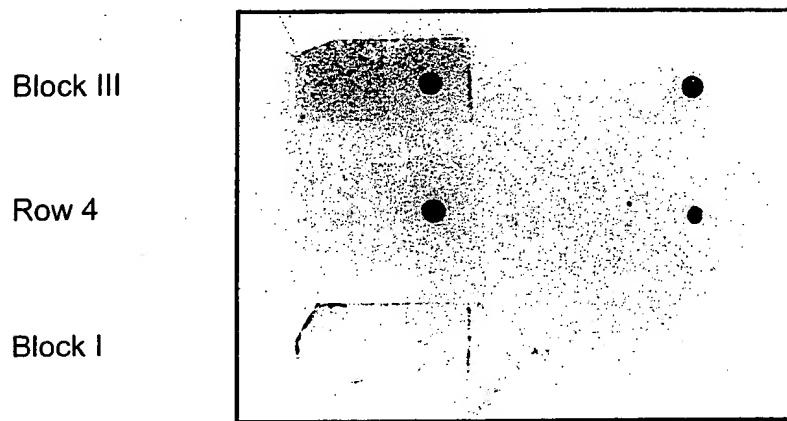


Figure 4b: Parallel screening for *Stig 1* mutant

49 plants (left side of the picture) or 100 plants (right side) were pooled and sampled. Two samples (Block III and Row 4) were known to contain an insertion mutant for the *Stig 1* gene, whereas one sample (Block I) was known not to contain this insert. Six replicate filters were prepared, harboring spotblotted DNA from the *Ap2A* gene (left side of each filter) or the *Stig 1* gene (right side).

The samples mentioned did not contain an insertion for the *Ap2A* gene. PCR products from each sample, obtained by Transposon Display, were fluorescein-labeled and used as a probe. Exposure time 5 minutes.